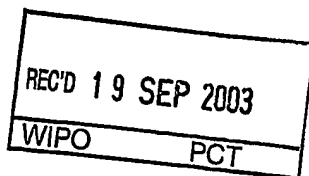




PCT/EP 03 / 08295



INVESTOR IN PEOPLE



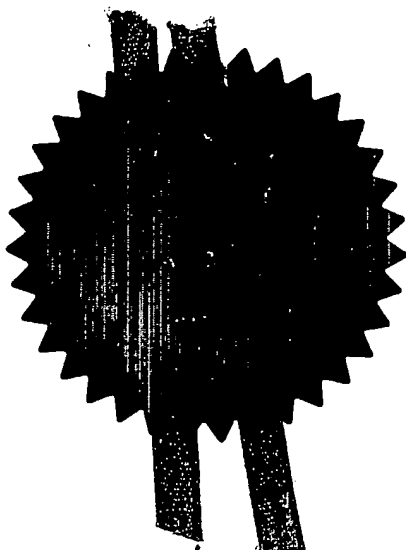
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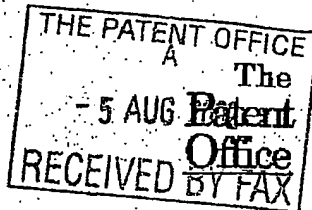
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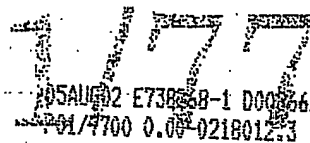
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PCT/EP 03 / 08295

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The Patent Office

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1. Your reference	GB Case	GT/3-22332/P1
2. Patent application number (The Patent Office will fill in this)	0218012.3 - 5 AUG 2002	
3. Full name, address and postcode of the or of each applicant (underline all surnames)	Ciba Specialty Chemicals Water Treatments Limited Cleckheaton Road Low Moor Bradford West Yorkshire BD12 0JZ	
Patent ADP number (if you know it)	7585391004	
If the applicant is a corporate body, give the country/state of its incorporation	England	
4. Title of invention	PRODUCTION OF A FERMENTATION PRODUCT	
5. Name of your agent (if you have one)	Ciba Specialty Chemicals Water Treatments Limited Patents Department PO Box 38 Cleckheaton Road Low Moor Bradford West Yorkshire BD12 0JZ	
"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)	7585391002	
6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and (if you know it) the or each application number	Country	Priority application number (if you know it)
		Date of filing (day/month/year)
7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application	Number of earlier application	Date of filing (day/month/year)
8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (Answer 'Yes' if:	YES	
a) any applicant named in part 3 is not an inventor, or		
b) there is an inventor who is not named as an applicant, or		
c) any named applicant is a corporate body.		
(see note (2))		

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Description 14

Claim(s) 4

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11.

I/We request the grant of a patent on the basis of this application

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5 August 2002

12. Name and daytime telephone number of person to contact in the United Kingdom

Catherine Choppen ..... 01274 417445

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1

DUPLICATE

### Production of a Fermentation Product

The present invention relates to processes of treating plant derived material to provide an aqueous liquor containing sugars which are used in a fermentation process to produce a fermentation product. Typically such fermentation products include for instance ethanol, glycerol, acetone, n-butanol, butanediol, isopropanol, butyric acid, methane, citric acid, fumaric acid, lactic acid, propionic acid, succinic acid, itaconic acid, acetic acid and acetaldehyde.

It is known to treat a biomass with acid in order to hydrolyse polysaccharides to the component sugars that can be used in a fermentation process to produce a fermentation product. For instance US-A-4384897 describes a method of treating biomass material in which it is subjected to a two stage hydrolysis in which polysaccharides that are more easily hydrolysed, such as hemicellulose and then in a second stage the more difficultly depolymerisable material e.g. cellulose, is depolymerised using a more severe hydrolytic treatment. The products of the first and second stages include sugar solutions and organic acids, aldehydes. The monosaccharides are subjected to fermentation to produce ethanol and the beer resulting from the fermentation may then be subjected to rectification to produce ethanol of commercial grade. US-A-4384897 sets out to provide improvements in more efficient washing of solids, the use of co-current washing or countercurrent washing of solids and proposes the use of ferric and or aluminium ions as flocculating agents to separate finely dispersed solids resulting from neutralisation of the hydrolysate liquor stream.

Kyoung Heon Kim et al (Applied Biochemistry and Biotechnology, Vol 91-93, pg 253-267) investigates the continuous countercurrent hydrolysis and extraction of hemicellulose from acid pretreated wood residues and considers the effect on drainage rate of such a pretreated biomass. A continuous countercurrent screw extractor used relies on the percolation of water by gravity through the pretreated biomass. One difficulty identified is that the pretreated biomass has poor water drainage properties and channelling or blockage may occur inside the extractor,

which can result in low sugar recovery or low throughput.

It would be desirable to improve the drainage properties of acid treated plant derived material in order to maximise sugar recovery.

According to the present invention we provide a process of producing fermentation product comprising the steps of,

- (i) forming an acidified suspension of particulate plant derived material comprising a first polysaccharide which is more readily hydrolysable and a second polysaccharide which is more difficult to hydrolyse,
- (ii) allowing the first polysaccharide to undergo hydrolysis by action of the acid at a temperature of at least 50°C under conditions such that the first polysaccharide is hydrolysed and thereby forming a mixture of an aqueous liquor containing dissolved sugar and a solid residue containing the second polysaccharide,
- (iii) subjecting the mixture to one or more separation stages in which the solid residue and aqueous sugar liquor are substantially separated from each other,
- (iv) optionally washing the residue substantially free of the sugar,
- (v) passing the solid cellulosic residue to a further treatment stage in which the residue is subjected to the action of dilute acid at a temperature of at least 50°C under conditions such that the second polysaccharide is hydrolysed and thereby forming a mixture of an aqueous liquor containing dissolved sugar and a solid residue,
- (vi) subjecting the mixture to one or more separation stages in which the solid residue and aqueous sugar liquor are substantially separated from each other,
- (vii) optionally washing the residue substantially free of the sugar,
- (viii) adjusting the pH of the aqueous liquor from stages (iii), (iv), (vi) and (vii) to a pH of at least 4,
- (ix) passing the aqueous liquor from stage (viii) to a fermentation stage in order to produce a fermentation product,

(x) separating the fermentation product from the broth, characterised in that the separation stage in steps (iii) and/or (vi) is assisted by flocculation of the by-product, employing one or more flocculating agent(s) selected from the group consisting of water soluble polymers, water swellable polymers and charged microparticulate material.

We have found that surprisingly by using the special flocculation process in the separation stage, a consistently high yield of fermentation product can be achieved. It is thought that small quantities of C<sub>5</sub> sugar resulting from the first stage remains with harder to hydrolyse material such as cellulose. Thus it would seem likely that the more severe hydrolysis conditions of the second stage hydrolysis results in these trace amounts of sugars to be converted into furfural and possibly other aldehydes. It is believed that even trace amounts of furfural and other aldehydes tend to poison the microorganisms or enzymic biocatalysts used in the fermentation process and thus resulting in the poor yields of fermentation product.

The improved separation stage in the process also has the advantage that the sugar solution resulting from the first and second stages is substantially free from extraneous solid material, such as cellulosic fibres.

The plant derived material is typically any readily available source of polysaccharides, particularly cellulosic materials. Typically the cellulosic material comprises materials selected from the group consisting of herbaceous biomass, softwood biomass, hardwood biomass, sewage sludge, paper mill sludge and the biomass fraction of municipal solid waste. The herbaceous biomass may for instance be bagasse, rice straw, rice hulls and cotton gin trash.

Preferably the plant derived material is cellulosic and comprises hemicellulose as the first polysaccharide and cellulose as the second polysaccharide. Generally the plant derived material also contains lignin or lignin type materials, which remain in the solid by-product.

The acidified suspension may be formed by combining a particulate material comprising cellulose, hemicellulose and lignin with a dilute acid. Alternatively the suspension can be made by treatment of a cellulosic biomass with sulphur dioxide gas, steam and water at an elevated temperature. Typically the process can be conducted by impregnation of the biomass material with SO<sub>2</sub> gas followed by steam at 205 to 215°C for 5 minutes and then the addition of water to form a slurry (Stenberg et al., 1998).

By dilute we mean that the acid generally has a concentration of less than 10% by weight. Usually though the concentration will be much lower, for instance at less than 5%. The acid may be a strong mineral acid such as hydrochloric acid, sulphuric acid, sulphurous acid, nitric acid or phosphoric acid. Suitable organic acids may be carbonic acid, tartaric acid, citric acid, glucuronic acid, formic acid, trichloro acetic acid or similar mono- or polycarboxylic acids.

Preferably the acid is a mixture of a strong mineral acid and an organic acid. Typically the mineral acid would be at a concentration of up to 2%, preferably in the range 0.2% to 1.0%, especially around 0.7%. The organic acid may be present at a higher concentration, for instance up to 5%, especially if the acid is a relatively weak acid such as acetic acid. Preferably the organic acid would be present at a concentration between 1% and 3%, more preferably around 2%.

The strong mineral ideally exhibits a pKa below 4. Preferred results are obtained by using either hydrochloric acid or sulphuric acid.

The hydrolysis of the first polysaccharide is preferably carried out at a temperature of between 120 and 220°C for a period of 1 to 15 minutes, although lower temperatures are possible if the treatment is longer. The hydrolysis of the second polysaccharide can be achieved at a temperature of between 120 and 220°C for a period of 1 to 15 minutes. Usually the second hydrolysis step is carried out at a higher temperature and/or for a longer period than the first hydrolysis step. However, given that the second polysaccharide is also subjected

to the first treatment stage, before removal of the hydrolysate resulting from the first polysaccharide, it may not always be necessary to subject the remaining second polysaccharide to an especially higher temperature and/or for longer and it may in fact be possible to achieve complete hydrolysis of the second polysaccharide at a lower temperature and/or for a shorter period than the first hydrolysis step.

In each of the first and second hydrolysis stages, the resulting hydrolysate is then separated from the solid materials, preferably through pressing of the treated material to separate the residue as a solid product. The solid product that is separated may be subjected to at least one wash cycle to remove any residual sugar solution from the solid. The wash cycle comprises washing the solid product with a suitable wash liquid. The wash liquid may be water. Normally the wash water is recycled water, for instance water that has been separated from the still bottoms liquor in the distillation recovery of the fermentation product in which suspended solids have been removed.

Since the wash water may contain other impurities which could be harmful to the fermentation process it would be desirable to minimise the amount of wash water used.

The liquid hydrolysate which contains sugars and acid can then be collected for further processing. When the first polysaccharide is hemicellulose, the resulting hydrolysate is generally C<sub>5</sub> sugars and when the second polysaccharide is cellulose the hydrolysate is generally C<sub>6</sub> sugars.

In each case it is important to adjust the pH of the acid sugar liquors to a pH of at least pH 4. The pH adjustment may be done by addition of a base or by use of an ion exchange resin, which is capable of absorbing the acid. Preferably the pH of the acidified aqueous sugar liquor that results from the digestion process is adjusted to a pH of at least 10 by addition of base material such as sodium carbonate, and then subsequent adjustment of the pH to more neutral or slightly acidic pH. Desirably the pH may be adjusted to a pH of between 10 and 12,



preferably about 11, by addition of a base, followed by titrating back to a pH of between 4 and 5, preferably about 4.5.

The acid may be removed from the liquor by adding the hydrolysate to a resin bed and the sugars are absorbed onto the resin and the sugars are absorbed onto the resin. The resin can then be purged with a gas substantially free of oxygen, which pushes the acid out of the resin. The resin can then be washed with water substantially free of oxygen. It would then be possible to produce an aqueous sugar stream. Ideally the sugar stream ideally contains at least 98% of the sugar present in the hydrolysate.

The C<sub>3</sub> and C<sub>4</sub> sugars may be fed into the fermentation vessel as separate streams or alternatively they may be combined into a single stream which is delivered into the fermentation vessel.

After the separation of the acid from the sugar stream, the acid is preferably concentrated for reuse. The concentration desirably can be achieved by evaporation.

The fermentation process of the present invention typically involves allowing the fermentation to proceed for 3 to 5 days. The fermentation product is separated from the broth by passing the broth comprising the fermentation product into a distillation stage, where the fermentation compound is collected as a distillate and the residue 'still bottoms' is removed. In one preferred aspect of the invention the fermentation product is separated from the broth by passing the broth comprising the fermentation product into a concentration stage, in which the fermentation compound is collected in the concentrate and extracted by at least one means selected from the group consisting of ion exchange, solvent extraction and electrodialysis.

The process can be used to prepare a range of fermentation products, but preferably the fermentation product is selected from the group consisting of

ethanol, glycerol, acetone, n-butanol, butanediol, isopropanol, butyric acid, methane, citric acid, fumaric acid, lactic acid, propionic acid, succinic acid, itaconic acid, acetic acid, acetaldehyde and 3-hydroxypropionic acid.

The microorganisms used in the fermentation process of the present invention can be, for example, a yeast such as *Kluyveromyces* species, *Candida* species, *Pichia* species, *Brettanomyces* species, *Saccharomyces* species such as *Saccharomyces cerevisiae* and *Saccharomyces uvarum*, *Hansenula* species and *Pachysolen* species. Alternatively, the microorganism can be a bacteria such as *Leuconostoc* species, *Enterobacter* species, *Klebsiella* species, *Erwinia* species, *Serratia* species, *Lactobacillus* species, *Lactococcus* species, *Pediococcus* species, *Clostridium* species, *Acetobacter* species, *Gluconobacter* species, *Lactobacillus* species, *Aspergillus* species, *Propionibacterium* species, *Rhizopus* species and *Zymomonas mobilis*. In addition genetically modified strains may also be used.

Since the solid product generally comprises lignin and analogous materials it can be particularly difficult to separate from the liquor. We have unexpectedly found that the production of fermentation product can be significantly improved by applying a particular flocculating agent to the separation of the hydrolysate from the solid product. We have found that the solid product can be more efficiently dewatered by the process and that a higher cake solids can be achieved. Since the solid product can be more efficiently dewatered there is a reduced requirement for separation equipment capacity and equipment that is less expensive and less expensive to operate, such as a filter press, can be used. Since higher cake solids can be achieved, less of the acid sugar solution remains in the residual by-product solid. Hence the quantity of water required to wash the by-product solid free of acid and sugar is much reduced.

Suitably the flocculating agent is selected from the group consisting of water soluble or water swellable natural, semi-natural and synthetic polymers.

Preferably the polymer is synthetic and may be formed by polymerisation of at

least one cationic, non-ionic or and/or anionic monomer(s) alone or with other water soluble monomers. By water soluble we mean that the monomer has a solubility of at least 5g/100 ml at 25°C.

Preferably polymeric flocculating agents are formed from ethylenically unsaturated water soluble monomers that readily polymerise to produce high molecular weight polymers. Particularly preferred polymers include monomers that are selected from the group consisting of polyacrylic acid, polyacrylamide, copolymers of acrylamide with (meth) acrylic acid or salts thereof, copolymers of acrylamide with dialkylaminoalkyl (meth) acrylate or acid addition or quaternary ammonium salts, polymers of diallyldimethyl ammonium chloride, polyamines and polyethylene imines. The polymers may be linear, branched or cross-linked.

The polymers may be prepared by any convenient conventional process, for instance by solution polymerisation, gel polymerisation, reverse phase suspension polymerisation and reverse phase emulsion polymerisation. Suitable processes include those described in EP-A-150933 or EP-A-102759.

Suitable polymers are anionic polymers. The preferred polymers are cationic polymers of sufficiently high molecular weight such that it exhibits an intrinsic viscosity of at least 4 dl/g. Such an intrinsic viscosity generally indicates a polymer of several million molecular weight, for instance generally greater than 5,000,000 and usually at least 7,000,000. In general the cationic polymer preferably has an intrinsic viscosity greater than 6 dl/g, often at least 8 or 9 dl/g. The intrinsic viscosity can be as high as 30 dl/g or higher. In many cases though suitable cationic polymers exhibit an intrinsic viscosity in the range of 7 to 25 dl/g, in particular 10 to 20 dl/g, in particular around 14 or 15 dl/g.

Suitable cationic monomers include quaternary ammonium or acid salts of monomers which contain amine groups. Preferably the cationic polymer is formed from a monomer or blend of monomers comprising at least one cationic monomer selected from the group consisting of quaternary ammonium and acid salts of

dimethylaminoethyl (meth) acrylate, quaternary ammonium and acid salts of dimethylaminoethyl (meth) acrylamide and diallyldimethyl ammonium chloride. The cationic monomers may be homopolymerised or copolymerised with other monomers, for instance acrylamide. The cationic polymers thus may be any polymer that carries a cationic, provided of course that they are of sufficiently high molecular weight to exhibit an intrinsic viscosity of at least 4 dl/g. Intrinsic viscosity is measured using a suspended level viscometer in 1M NaCl buffered to pH 7.5 at 25°C.

The cationic polymers according to the invention may be prepared as substantially linear polymers or as branched or structured polymers. Structured or branched polymers are usually prepared by inclusion of polyethylenically unsaturated monomers, such as methylene-bis-acrylamide into the monomer mix, for instance as given in EP-B-202780. Preferably however, the polymers are substantially linear and are prepared in the form of a bead or powdered product.

Suitably the polymeric flocculating agent would be added as an aqueous solution or aqueous dispersion. The polymer may be added in an amount sufficient to effect flocculation. Typically the amount of polymeric flocculating agent sufficient to induce flocculation would be usually at least 0.002 weight % based on weight of suspended solids. Usually better flocculation and therefore separation can be achieved if at least 0.01% is used. The dose may be substantially higher, for instance up to 1%. However, optimum flocculation and separation is normally achieved using doses in the range of 0.015% to 0.2%, especially 0.02% to 0.1%. Following flocculation of the suspended solids the solid product can be separated from the hydrolysate aqueous liquor by mechanical means, for instance filter press, centrifuge, belt press, horizontal belt filter or pressure filter. The action of the flocculating agent greatly enhances the separation of the solids from the liquor by comparison to separation using solely mechanical means. We have found that the process of the present invention provides a higher cake solids, with less residual aqueous liquor, which means that a higher proportion of the sugar liquor is available for conversion into the fermentation product. Likewise we find that the

aqueous liquor contains much lower levels of extraneous suspended cellulosic solids. Furthermore we also find that less wash water is required.

The solid product that results from the separation step should be as dry as possible in order to prevent any loss of sugar, which would otherwise be used in the fermentation process. In addition the solid by product, which contains mainly lignin can be used as a solid fuel, for instance for use in the boiler used to heat the distillation column used in separating the fermentation product from the fermentation broth. Thus it is also important that when the solid lignin containing product is used as a fuel that it is as dry as possible.

In a further preferred embodiment of the present invention the flocculating agent is a charged microparticulate material. Particularly suitable examples of charged microparticulate materials include swellable clays, anionic, cationic or amphoteric microparticulate silica based materials and organic cross-linked polymeric microparticles.

The siliceous material may be any of the materials selected from the group consisting of silica based particles, silica microgels, colloidal silica, silica sols, silica gels, polysilicates, aluminosilicates, polyaluminosilicates, borosilicates, polyborosilicates, zeolites or swellable clay.

This siliceous material may be in the form of an anionic microparticulate material. Alternatively the siliceous material may be a cationic silica. Desirably the siliceous material may be selected from silicas and polysilicates.

The polysilicates of the invention may be prepared by acidifying an aqueous solution of an alkali metal silicate. For instance polysilicic microgels otherwise known as active silica may be prepared by acidification of alkali metal silicate to between pH 2 and 10 by use of mineral acids or acid exchange resins, acid salts and acid gases. It may be desired to age the freshly formed polysilicic acid in order to allow sufficient three dimensional network structure to form. Generally the

time of ageing is insufficient for the polysilicic acid to gel. Particularly preferred siliceous material include polyalumino-silicates. The polyaluminosilicates may be for instance aluminated polysilicic acid, made by first forming polysilicic acid microparticles and then post treating with aluminium salts.

Alternatively the polyaluminosilicates may be polyparticulate polysilicic microgels of surface area in excess of  $1000\text{m}^2/\text{g}$  formed by reacting an alkali metal silicate with acid and water soluble aluminium salts. Typically the polyaluminosilicates may have a mole ratio of alumina:silica of between 1:10 and 1:1500.

Polyaluminosilicates may be formed by acidifying an aqueous solution of alkali metal silicate to between pH 2 and 10 using concentrated sulphuric acid containing 0.2 to 2.0% by weight of a water soluble aluminium salt, for instance aluminium sulphate. The aqueous solution may be aged sufficiently for the three dimensional microgel to form. Typically the polyaluminosilicate is aged for up to about two and a half hours before diluting the aqueous polysilicate to 0.5 weight % of silica.

The siliceous material may be a colloidal borosilicate. The colloidal borosilicate may be prepared by contacting a dilute aqueous solution of an alkali metal silicate with a cation exchange resin to produce a silicic acid and then forming a gel by mixing together a dilute aqueous solution of an alkali metal borate with an alkali metal hydroxide to form an aqueous solution containing 0.01 to 30 %  $\text{B}_2\text{O}_3$ , having a pH of from 7 to 10.5.

The swellable clays may for instance be typically a bentonite type clay. The preferred clays are swellable in water and include clays which are naturally water swellable or clays which can be modified, for instance by ion exchange to render them water swellable. Suitable water swellable clays include but are not limited to clays often referred to as hectorite, smectites, montmorillonites, nontronites, saponite, sauconite, hornites, attapulgites and sepiolites.

Most preferably the clay is a bentonite type clay. The bentonite may be provided as an alkali metal bentonite. Bentonites occur naturally either as alkaline

bentonites, such as sodium bentonite or as the alkaline earth metal salt, usually the calcium or magnesium salt. Generally the alkaline earth metal bentonites are activated by treatment with sodium carbonate or sodium bicarbonate. Activated swellable bentonite clay is often supplied to the paper mill as dry powder. Alternatively the bentonite may be provided as a high solids flowable slurry, for example at least 15 or 20% solids.

When the charged microparticulate material comprises an organic cross-linked polymeric microparticles. The microparticles may be made as microemulsions by a process employing an aqueous solution comprising a cationic or anionic monomer and crosslinking agent; an oil comprising a saturated hydrocarbon; and an effective amount of a surfactant sufficient to produce particles of less than about 0.75 micron in unswollen number average particle size diameter.

Microbeads are also made as microgels by procedures described by Ying Huang et. al., Makromol. Chem. 186, 273-281 (1985) or may be obtained commercially as microlatices. The term "microparticle", as used herein, is meant to include all of these configurations, i.e. beads per se, microgels and microlatices.

The charged microparticle material may be used in amounts of at least 0.002% based on weight of suspended solids. Typically though the doses are usually as high as 0.8 or 1.0% or higher. When the charged microparticle material is inorganic, the dose is usually in excess of 0.06%, preferably in the range 0.1 to 0.6%. When the charged microparticle is organic the dose is typically below 0.3%, preferably in the range 0.02 to 0.1%.

Unexpectedly we have found that the hydrolysate liquor can be separated particularly rapidly when the flocculation is effected by employing a water soluble or water-swellable polymer and a charged microparticulate material. In one aspect we find that particularly effective flocculation and separation of the solids from the liquor is effected when flocculation is effected by introducing an anionic microparticle material into the mixture and then reflocculating by adding a cationic or substantially non-ionic polymer. In a further preferred embodiment of the

present invention we find that especially fast and efficient separation of solids is achieved by a process in which flocculation is effected by introducing a cationic polymer into the mixture and then reflocculating by adding an anionic microparticulate material.

The following example illustrates the invention.

#### Example

100g of softwood chips ground to less than 2mm were added to 400 g of water and heated to 190°C. Once at 190°C sulphuric acid was added to a concentration of 0.7% under nitrogen pressure and the mixture was left for 3 minutes. The temperature was rapidly reduced to 80°C and the insoluble solids present in the pre-hydrolysed slurry (containing 0.32% sulphuric acid) is then separated on a filter press. Solutions of flocculant or flocculants (at 0.2 to 0.5% solids) and/or microparticulate suspensions (at 0.5 to 15% solids) are added into the slurry with necessary agitation at a dose of 0.2 to 2 Kg per tonne of solids. The flocculants were found to increase the rate of free drainage by gravity through a porous belt, before preparation of a filter cake in a wedge zone and subsequent further dewatering in a pressure zone.

A method of removing the sugars remaining in the solid portion is to wash with (recycled) water.

The recovered solid portion is again prepared as a slurry with approximately 400 ml of water and heated to 215°C. Once at 215°C sulphuric acid is added to a concentration of 0.4% under nitrogen and the mixture was left for 3 minutes. The temperature was rapidly reduced to 80°C and the insoluble solids present in the pre-hydrolysed slurry (containing 0.32% sulphuric acid) is then separated on a filter press. Prior to pressing solutions of flocculant or flocculants (at 0.2 to 0.5% solids) and/or particulate suspensions (at 0.5 to 15% solids) are added into the slurry with necessary agitation at a dose of 0.2 to 2 Kg per tonne of solids.

Flocculants increase the rate

of free drainage by gravity through a porous belt, before preparation of a filter



cake in a wedge zone and subsequent further dewatering in a pressure zone. The flocculated dewatered pressure belt filter cake containing predominantly lignin was further dewatered in a pressure filter to generate a cake with a high dry solids concentration (approximately 85%(w/w)) suitable for use as a fuel. A method of removing the sugars remaining in the solid portion is to wash with (recycled) water.

After ion exchange for the removal of acetic acid, the liquid portion of the hydrolyzate is acidified to pH 2 by the addition of sulphuric acid. Lime is then added to raise the pH to 10 and heated to 50°C. The liquid is then adjusted to the fermentation pH of 4.5 for 4 hours allowing gypsum crystals to form for separation by filtration.

Following the two-stage dilute acid hydrolysis the solid residues remaining can either enter the fermentation process or be separated for fuel.

**Claims**

1. A process of producing fermentation product comprising the steps of,
- (i) forming an acidified suspension of particulate plant derived material comprising a first polysaccharide which is more readily hydrolysable and a second polysaccharide which is more difficult to hydrolyse,
  - (ii) allowing the first polysaccharide to undergo hydrolysis by action of the acid at a temperature of at least 50°C under conditions such that the first polysaccharide is hydrolysed and thereby forming a mixture of an aqueous liquor containing dissolved sugar and a solid residue containing the second polysaccharide,
  - (iii) subjecting the mixture to one or more separation stages in which the solid residue and aqueous sugar liquor are substantially separated from each other,
  - (iv) optionally washing the residue substantially free of the sugar,
  - (v) passing the solid cellulosic residue to a further treatment stage in which the residue is subjected to the action of dilute acid at a temperature of at least 50°C under conditions such that the second polysaccharide is hydrolysed and thereby forming a mixture of an aqueous liquor containing dissolved sugar and a solid residue,
  - (vi) subjecting the mixture to one or more separation stages in which the solid residue and aqueous sugar liquor are substantially separated from each other,
  - (vii) optionally washing the residue substantially free of the sugar,
  - (viii) adjusting the pH of the aqueous liquor from stages (iii), (iv), (vi) and (vii) to a pH of at least 4,
  - (ix) passing the aqueous liquor from stage (viii) to a fermentation stage where the dissolved sugars are acted upon by an enzyme in a fermentation broth to produce a fermentation product,
  - (x) separating the fermentation product from the broth,

characterised in that the separation stage in steps (iii) and/or (vi) is assisted by flocculation of the waste by-product, employing one or more flocculating agent(s) selected from the group consisting of water soluble polymers, water swellable

polymers and charged microparticulate material.

2. A process according to claim 1 in which the plant derived material comprises materials selected from the group consisting of herbaceous biomass, softwood biomass, hardwood biomass, sewage sludge, paper mill sludge and the biomass fraction of municipal solid waste.

3. A process according to claim 1 or claim 2 in which the plant derived material is cellulosic and comprises hemicellulose as the first polysaccharide and cellulose as the second polysaccharide.

4. A process according to any of claims 1 to 3 in which the acid has a pKa of below 4 and has a concentration up to 2% by weight.

5. A process according any of claims 1 to 4 in which the acid is selected from sulphuric acid and hydrochloric acid.

6. A process according to any of claims 1 to 5 in which the hydrolysis of the first polysaccharide is conducted at a temperature of between 120 to 220°C for a period of from 1 minute to 15 minutes.

7. A process according to any of claims 1 to 6 in which the hydrolysis of the second polysaccharide is conducted at a temperature of between 120 to 220°C for a period of from 1 minute to 15 minutes.

8. A process according to any of claims 1 to 7 in which the flocculating agent is selected from the group consisting of water soluble or water swellable natural, semi-natural and synthetic polymers.

9. A process according to claim 8 in which the polymer is formed from a water soluble monomer or blend of monomers.

10. A process according to claim 8 in which the polymer is selected from the group consisting of polyacrylate salts, polyacrylamide, copolymers of acrylamide with (meth) acrylic acid or salts thereof, copolymers of acrylamide with dialkylaminoalkyl (meth) acrylate or acid addition or quaternary ammonium salts, polymers of diallyldimethyl ammonium chloride, polyamines and polyethylene imines.

11. A process according to any of claims 1 to 10 in which the flocculating agent is a charged microparticulate material.

12. A process according claim 11 in which the charged microparticulate

material is selected from the group consisting of swellable clays, anionic, cationic or amphoteric microparticulate silica based materials and organic cross-linked polymeric microparticles.

13. A process according to any one of claims 1 to 12 in which flocculation is effected by employing a water soluble or water-swellable polymer and a charged microparticulate material.

14. A process according to any one of claims 1 to 13 in which flocculation is effected by introducing an anionic microparticle material into the mixture and then reflocculating by adding a substantially non-ionic polymer.

15. A process according to any one of claims 1 to 14 in which flocculation is effected by introducing a cationic polymer into the mixture and then reflocculating by adding an anionic microparticulate material.

16. A process according to any one of claims 1 to 15 in which flocculation is effected by introducing a cationic polymer into the mixture and then reflocculating by adding an anionic polymer.

17. A process according to any one of claims 1 to 16 in which flocculation is effected by introducing an anionic polymer into the mixture and then reflocculating by adding a cationic polymer.

18. A process according to any of claims 1 to 17 in which the solid-by product material comprises lignin.

19. A process according to any of claims 1 to 18 in which the fermentation product is selected from the group consisting of ethanol, glycerol, acetone, n-butanol, butanediol, isopropanol, butyric acid, methane, citric acid, fumaric acid, lactic acid, propionic acid, succinic acid, itaconic acid, acetic acid, acetaldehyde and 3-hydroxypropionic acid.

20. A process according to any of claims 1 to 19 in which the fermentation product is separated from the broth by passing the broth comprising the fermentation product into a distillation stage, where the fermentation compound is collected as a distillate and the residue 'still bottoms' is removed.

21. A process according to any one of claims 1 to 20 in which the fermentation product is separated from the broth by passing the broth comprising the fermentation product into a concentration stage, in which the fermentation

compound is collected in the concentrate and extracted by at least one means selected from the group consisting of ion exchange, solvent extraction and electro dialysis.

**Abstract**

A process of producing fermentation product comprising the steps of,

- (i) forming an acidified suspension of particulate plant derived material comprising a first polysaccharide which is more readily hydrolysable and a second polysaccharide which is more difficult to hydrolyse,
- (ii) allowing the first polysaccharide to undergo hydrolysis by action of the acid at a temperature of at least 50°C under conditions such that the first polysaccharide is hydrolysed and thereby forming a mixture of an aqueous liquor containing dissolved sugar and a solid residue containing the second polysaccharide,
- (iii) subjecting the mixture to one or more separation stages in which the solid residue and aqueous sugar liquor are substantially separated from each other,
- (iv) optionally washing the residue substantially free of the sugar,
- (v) passing the solid cellulosic residue to a further treatment stage in which the residue is subjected to the action of dilute acid at a temperature of at least 50°C under conditions such that the second polysaccharide is hydrolysed and thereby forming a mixture of an aqueous liquor containing dissolved sugar and a solid residue,
- (vi) subjecting the mixture to one or more separation stages in which the solid residue and aqueous sugar liquor are substantially separated from each other,
- (vii) optionally washing the residue substantially free of the sugar,
- (viii) adjusting the pH of the aqueous liquor from stages (iii), (iv), (vi) and (vii) to a pH of at least 4,
- (ix) passing the aqueous liquor from stage (viii) to a fermentation stage in order to produce a fermentation product,
- (x) separating the fermentation product from the broth,

wherein the separation stage in steps (iii) and/or (vi) is assisted by flocculation of the waste by-product, employing one or more flocculating agent(s) which are water soluble or swellable polymers or charged microparticulate material.

Typically such fermentation products include for instance ethanol, glycerol,

acetone, n-butanol, butanediol, isopropanol, butyric acid, methane, citric acid,  
fumaric acid, lactic acid, propionic acid, succinic acid, itaconic acid, acetic acid,  
acetaldehyde and 3-hydroxypropionic acid.

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